Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

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- 1. (Withdrawn) A method for treatment of chronic pain comprising orally administering the composition of claim 9.
- 2. (Withdrawn) The method of claim 1 wherein said tricyclic antidepressant is administered in a dosage of from about 2.5 mg to about 25 mg daily.
- 3. (Withdrawn) The method of claim 2 wherein said tricyclic antidepressant compound is selected from the group consisting of doxepin, amitriptyline, desipramine, imipramine and physiologically acceptable acid addition salts thereof.
- 4. (Withdrawn) The method of claim wherein said physiologically acceptable acid addition salts are selected from the group consisting of the hydrochloride, hydrobromide, hydroiodide, acetate, valerate andoleate.
- 5. (Withdrawn) The method of claim 1 wherein said non-narcotic analysesic is administered in a dosage from about 0.50 gms to about 2.6 gms daily.
- 7. (Withdrawn) T he method of claim 2 wherein said low dose of tricyclic antidepressant compound and said standard dose of non-narcotic analgesic are present in a single composition including a pharmaceutically acceptable vehicle for oral administration.
- 8. (Withdrawn) The method of claim 7 wherein said composition is in a form selected from the group consisting of tablets, capsules, caplets, oral solutions, and oral suspensions.
- 9. (Previously presented) A composition for treatment of chronic pain consisting essentially of a combination of a low dose of a tricyclic antidepressant compound and a standard dose of a non-narcotic analysis in a pharmaceutical acceptable vehicle for oral administration.
- 10. (Previously presented) The composition of claim 9 being provided in a daily dosage form wherein said tricyclic antidepressant compound is present in an amount of about 2.5 mg to 25 mg.
- 11. (Original) The composition of claim 9 wherein said tricyclic antidepressant compound is selected from the group consisting of doxepin, amitriptyline, desipramine, imipramine, and physiologically acceptable acid addition salts thereof.

- 12. (Original) The composition of claim 9 wherein said physiologically acceptable acid addition salts are selected from the group consisting of the hydrochloride, hydrobromide, hydroiodide, acetate, valerate and oleate.
- 13. (Previously presented) The composition of claim 9 wherein said non-narcotic analgesic is administered in a dosage of from about 0.50 gms to about 2.6 gms daily.
- 14. (Previously presented) The composition of claim 9 wherein said non-narcotic analgesic is selected from the group consisting of acetaminophen or non-steroidal anti-inflammatory drugs.
- 15. (Original) The composition of claim 7 wherein the combination of a tricyclic antidepressant and a non-narcotic analgesic and a pharmaceutically acceptable vehicle is in a form selected from the group consisting of tablets, capsules, caplets, oral solutions and oral suspensions.